

Q4 and FY 2018 Results

Media Presentation January 30, 2019



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Agenda

1. Company overview

Vas Narasimhan – CEO Novartis

2. Financial review

Harry Kirsch – CFO Novartis

3. Q&A

Novartis Executive Committee members

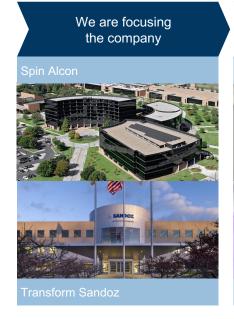


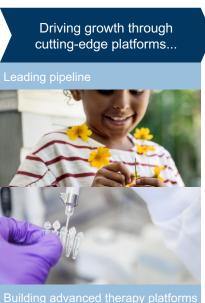
Company overview



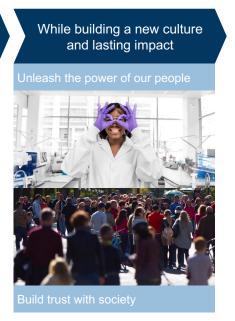
We aim to become a leading medicines company

Powered by advanced therapy platforms and data science









We delivered strong growth with operating leverage in Q4 and FY 2018

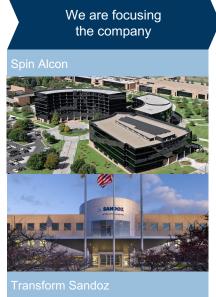
	Q4 (% cc)	Full year (% cc)		
	Sales	Sales Core OpInc		Core OpInc	
Group ¹	+6% ▲	+11% 🔺	+5%	+8% 🛕	
Innovative Medicines	+9%	+13%	+8%	+11%	
Sandoz	-2%	-5%	-3%	-3%	
Alcon	+4%	0%	+5%	+10%	

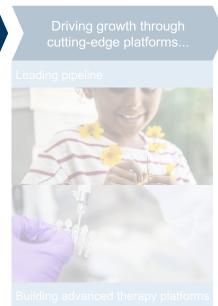
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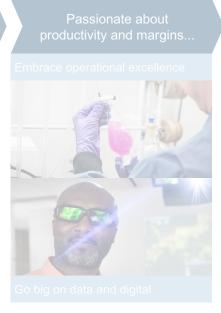


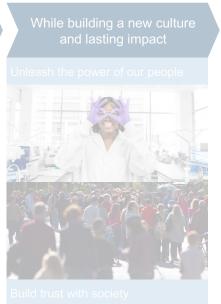
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We took major steps to focus the company in 2018, while building leading advanced therapy platforms

Exits¹ to focus the company Deals¹ to build new platforms Spark Consumer Healthcare Alcon² Gene therapy Cell therapy Radioligand therapy

All trademarks are the property of their respective owners 1. Announced or closed in 2018 2. The planned 100% spin-off of Alcon remains subject to certain conditions precedent, such as no material adverse events, receipt of necessary authorizations as well as tax rulings and opinions and shareholder approval at the AGM in February 2019; completion expected in H1 2019 3. The announced sale of Sandoz US dermatology and oral solids portfolio to Aurobindo is subject to the completion of customary closing conditions expected in 2019 4. Sale of our anti-bacterial portfolio to Boston Pharmaceuticals is one example of portfolio prioritization. Others include out-licensing of BJG398 to QED Therapeutics, FGF401 to EverNov. LXS196 to IDEAYA.



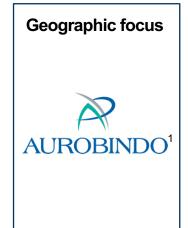
pharmaceuticals

Our Sandoz transformation will help enable us to compete in a more challenging environment

Reshaping the portfolio...



...while driving efficiency



Lean cost structure

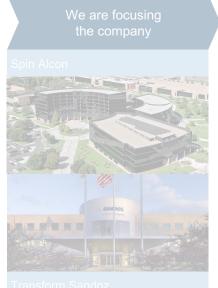
- SKU rationalization
- Manufacturing footprint optimization
- Regional consolidation

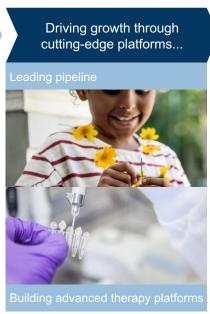
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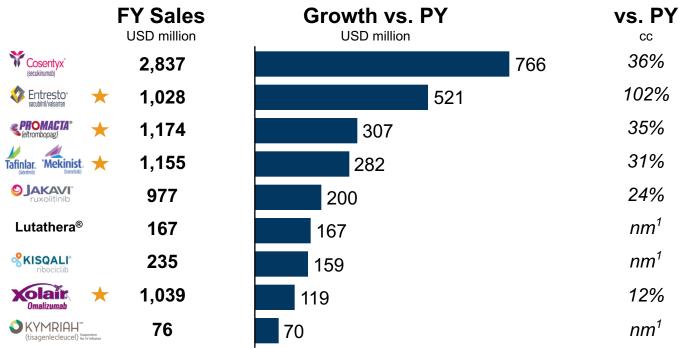






Our in-line growth brands provide a solid foundation for continued growth

★ 4 additional blockbusters in 2018



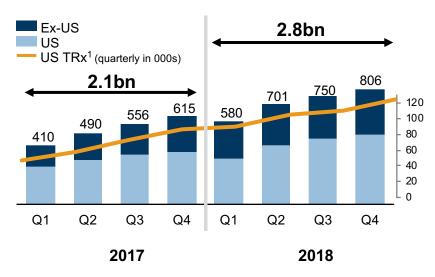
¹ Not meaningful



Cosentyx®: Strong growth driven by demand, well positioned across indications

Quarterly sales evolution

USD million



1. IMS NPA TRx, Cosentyx® restated in Aug 2017 to include free product, Q4 estimated with WE 12/21/2018 data

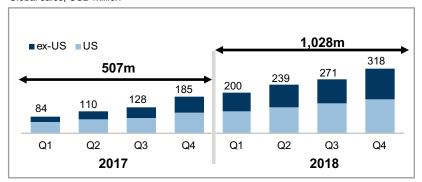
- Q4 sales USD 806m (+33% cc) with consistent growth US (+34% cc) and ex-US (+32% cc)
 - Continued demand growth, US YoY TRx +29% Dermatology, +49% in Rheumatology¹
 - Gaining share in a growing and competitive psoriasis market
- 2019: expected to maintain strong access in US
- Continue to advance science in psoriatic disease
 - ARROW (readout expected end of 2019) expected to confirm importance of IL-17A vs. IL-23
 - Cosentyx® has demonstrated efficacy in the multiple manifestations of psoriatic disease
- PREVENT (nrAxSpA) read-out and submission expected end 2019



Entresto[®] achieves blockbuster status – reinforcing strong therapy position in heart failure¹

Strong sales growth driven by execution & new data

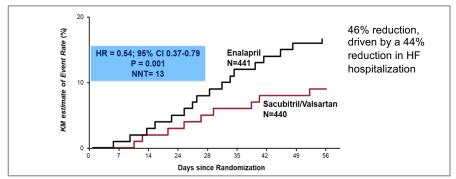
Global sales, USD million



- USD 318m (+76% cc) Q4 sales
- Blockbuster in 2018 and doubling sales vs. 2017

PIONEER data² supports early Entresto® use

Composite of Death, HF re-hospitalization, LVAD, Listing for Transplant



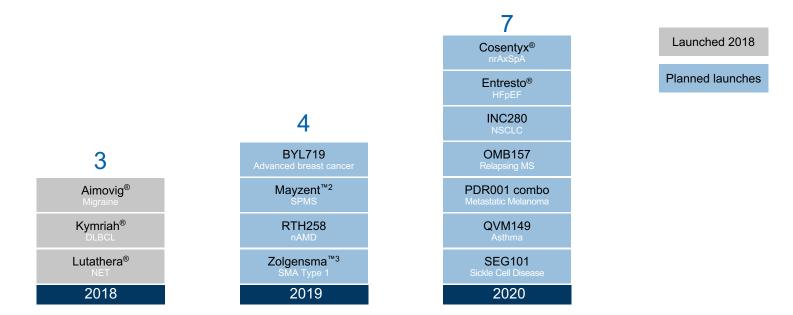
- Early indicators of accelerated hospital initiation (PIONEER & TRANSITION)
- Significantly reduces NT-proBNP in stabilized ADHF patients (PIONEER)

Expected newsflow 2019 FIR: PARALLEL-HF (Japan registration trial HFrEF) Q2 2019; PARAGON-HF in HFpEF Q3 2019

ADHF – Acute Decompensated Heart Failure FIR – First Interpretable Results LVAD – Left Ventricular Assist Device 1. Entresto® is approved in HFrEF and ongoing HFpEF trial expected to read out 2019. 2. Published in New England Journal of Medicine. Nov 2018; DOI: 10.1056/NEJMoa1812851



With 10+ potential blockbuster¹ launches planned in the next 2 years



^{1.} Individual assets with expected peak sales >USD 1bn across all indications 2. The brand name Mayzent™ has been provisionally approved by the FDA and EMA for the investigational product siponimod (BAF312), but the product itself has not been approved for sale in any country 3. The brand name Zolgensma™ has been provisionally approved by the FDA for the investigational product AVXS-101 (onasemnogene abeparvovec-xxxx), but the product itself has not received marketing authorization or BLA approval from any regulatory authorities



2019 expected catalysts to continue the momentum

Catalysts		Selected examples			
Key approvals	15	Zolgensma™ ¹ SMA Type 1 (US/EU/JP)	Brolucizumab (RTH258) Neovascular AMD (US)		
аррготаю		Mayzent™² SPMS (US/EU/JP)	Alpelisib (BYL719) Breast Cancer (US)		
Major submissions	20	Ofatumumab (OMB157) Relapsing MS (US/EU)	Brolucizumab (RTH258) PDR001 combo Neovascular AMD (US/EU/JP) Metastatic Melanoma (US/		
		Crizanlizumab (SEG101) Sickle Cell Disease (US/EU)	INC280 NSCLC (US/JP)		
Major late-stage readouts	6	Zolgensma^{™1} SMA Type 2	Entresto [®] HFpEF	Ofatumumab (OMB157) Relapsing MS	
		Fevipiprant (QAW039) Asthma	Cosentyx ® nrAxSpA	PDR001 combo Metastatic Melanoma	

^{1.} The brand name Zolgensma™ has been provisionally approved by the FDA for the investigational product AVXS-101 (onasemnogene abeparvovec-xxxx), but the product itself has not received marketing authorization or BLA approval from any regulatory authorities 2. The brand name MayzentTM has been provisionally approved by the FDA and EMA for the investigational product siponimod (BAF312), but the product itself has not been approved for sale in any country



Advanced therapy platforms with the potential to expand the game-board

(Illus	strative)	Small molecules	Large molecules	Cell therapy	Gene therapy	Radioligand therapy
مگی	Oncology		Bispecific antibodies ²	CAR-T CBMG ⁵		AAA Endocyte
U	Cardio-Metabolic		materials ¹	Intellia & Cell for Cure ⁴		
	IHD	degradation				
R	Neuroscience	Transcription factors			NIBR Portfolio	
	Ophthalmology		Novel bio-materials		AveXis Luxturna®	
A	Respiratory		Inhaled biologics			

^{1.} Partnership with the Wyss Institute for Biologically Inspired Engineering at Harvard University and the Dana-Farber Cancer Institute 2. Collaboration with Xencor 3. Collaborations with Intellia Therapeutics and Caribou Biosciences 4. Proposed acquisition; transaction subject to the completion of customary closing conditions 5. Collaboration with Cellular Biomedicine Group in China



Gene therapy platform with rapidly expanding pipeline, deep manufacturing expertise in AAV



7 programs in clinic over next year

Selected assets	Indication	Stage	Next milestone
AVXS-101 (AAV9)	SMA	Filed	Regulatory approval(s) 1H19
CGF166 (Ad5)	Hearing loss	Phase 1	
CPK850 (AAV8)	Retinitis pigmentosa	Phase 1	
AVXS-201 RTT (AAV9)	Rett Syndrome	Preclinical	IND 1Q19
AVXS-301 SOD1 (AAV9)	Inherited ALS-SOD1	Preclinical	IND 2Q19
AVXS-401	Undisclosed	Preclinical	IND 2H19
AVXS-501	Undisclosed	Preclinical	IND 4Q19 / 1Q20

Manufacturing expansion ongoing for AAV-based gene therapies



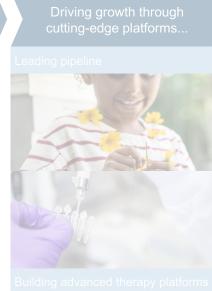
- Flexible, disposable manufacturing platform
- Operational facility in Chicago; ongoing buildout in North Carolina
- Capabilities across AAV vectors; deep CMC expertise



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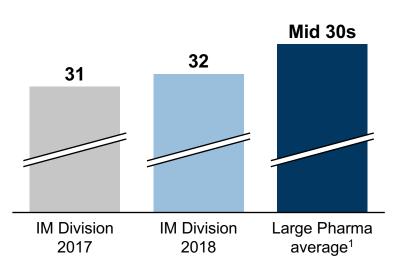




We are committed to driving consistent margin expansion

Innovative Medicines

Core margin (%)



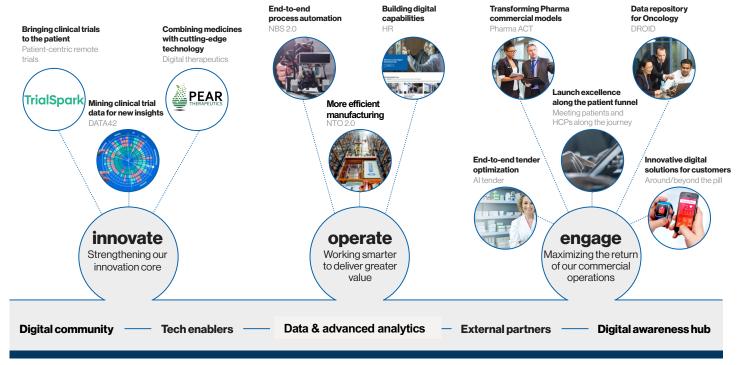
- Resource allocation and productivity programs in commercial units
- Cross-divisional synergies: Novartis Technical Operations, Novartis Business Services, Procurement
- Generics (mainly Afinitor[®], Exjade[®], Gilenya[®])
- Launch investments for potential future blockbusters



Acceleration of key growth drivers

^{1.} Source: Novartis analysis of average 2016 core margin of Large Pharma peer companies

Advancing an enterprise-wide digital transformation

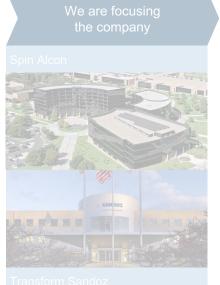


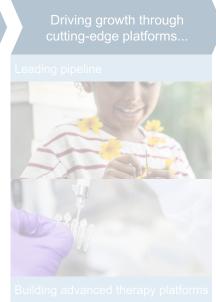
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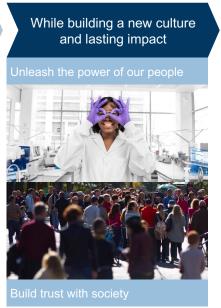
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Culture transformation is key to our success

Initiated 5-year journey in 2018





Focused effort to build lasting trust with society



Ethical **Standards**

Established Ethics, Risk & Compliance function, led by **Executive Committee member**

Embedding principles-based decision-making in the organization



Pricing and Access

Integrating Access Principles into overall business strategy

Improved ranking in Access to Medicines Index to #2



Global Health Challenges

Renewed commitment to malaria with USD 100m investment

Established Global Partnership for Zero Leprosy



Corporate Citizenship

Approved new environmental sustainability targets, incl. carbon neutrality by 2025

Helped lead Pat-INFORMED initiative, making patents available online



Stakeholder Engagement

Continued to improve transparency and evolve reporting

Increased reporting on Financial, Environmental and Social (FES) impact on society



Financial review

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2018 financial results in line with guidance

Group full year guidance (January 2018)
In cc

"Sales are expected to grow low to mid single digit"

"Core operating income expected to grow mid to high single digit"

8%

✓



Summary of Q4 and FY 2018 financial results

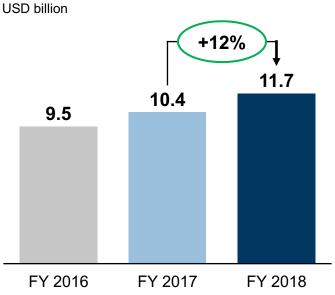
Group ¹	Q4	Change vs. PY		FY	Change vs. PY	
USD million	2018	% USD	% сс	2018	% USD	% сс
Net Sales	13,269	3	6	51,900	6	5
Core Operating income	3,387	5	11)	13,823	8	8
Operating income	1,299	-37	-29	8,169	-5	-5
Net Income	1,194	-40	-32	12,614	64	64
Core EPS (USD)	1.25	3	9	5.15	6	6
EPS (USD)	0.52	-39	-32	5.44	66	66
Free Cash Flow	2,939	20		11,717	12	

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FY 2018 free cash flow at USD 11.7bn

Group free cash flow

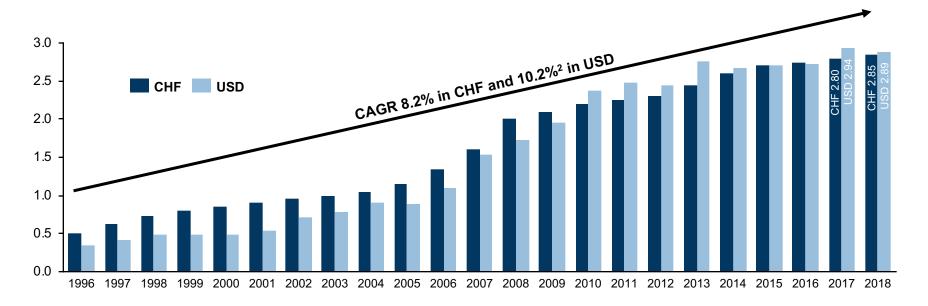


2018 Key drivers vs. PY:

- Cash flows from operating activities, mainly:
 - + Higher core operating income
 - + GSK milestone receipt (divested Vaccines business)
- Higher net intangible investments



Novartis proposes 22nd consecutive dividend increase to the AGM: 2.85 CHF / share¹



^{1.} Proposal to shareholders at the 2019 Annual General Meeting, taking place on February 28, 2019 2. Converted at historic exchange rates at the dividend payment dates as per Bloomberg; assumes an exchange rate of USD / CHF of 0.9862 as of December 31, 2018 for 2018



2019 Novartis full year guidance

Barring unforeseen events (in cc); Growth vs PY in cc

Cu	rrent	t Gro u	ıp stru	ıcture

No change to current Group Structure

Net Sales

Expected to grow low to mid single digit

- IM Division to grow mid single digit
- Sandoz to decline low single digit
- Alcon to grow low to mid single digit

Core Operating Income

Expected to grow mid single digit

Alcon Core OpInc margin expected to expand

New focused medicines company

Excl. Alcon¹ & Sandoz proposed US portfolio sale to Aurobindo² from both 2018 and 2019

Expected to grow mid single digit

- IM Division to arow mid single digit
- Sandoz to be broadly in line with prior year

Expected to grow mid to high single digit

Key Assumption: All guidance includes forecast assumption that no Gilenya® generics enter in 2019. However, generic competitors may still launch at risk

^{1.} The planned 100% spinoff of Alcon remains subject to certain conditions precedent, such as no material adverse events, receipt of necessary authorizations as well as tax rulings and opinions and shareholder approval at the AGM in February 2019; completion expected in H1 2019 2. The announced sale of Sandoz US dermatology and oral solids portfolio to Aurobindo, expected to close during 2019, is subject to the completion of customary closing conditions. Estimated 2018 FY Sales and Core Opinc of the Sandoz US Oral solids and Dermatology businesses were approximately USD 1.2bn and 0.3bn, respectively.



Closing

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We will continue to drive our priorities in 2019



- Deliver pipeline targets, incl. Zolgensma™, Mayzent™, RTH258 and BYL719 approvals
- Maintain high proportion of first-in-class / transformative assets
- Extend leadership in cell, gene and radioligand therapies



Operational Excellence

- Execute Alcon spin and progress Sandoz transformation
- Drive productivity through NTO and NBS transformations
- Deliver performance of in-market growth drivers
- Prepare for 10+ potential blockbuster launches, incl. 4 in 2019



Data & Digital Leadership

- Scale top 5 digital initiatives across the company
- Upskill digital capabilities in all units and functions
- Build Novartis position within digital ecosystem



- Continue to embed principles-based decision-making (P3)
- Implement Novartis Access Principles, with every new innovative medicine launch having an access plan
- Progress efforts in global health



Culture

- Continue 5-year journey to transform culture, with a focus on strengthening leadership capabilities
- Further increase diversity and inclusion

Deliver financials

Sales growth and margin expansion



A&Q

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